

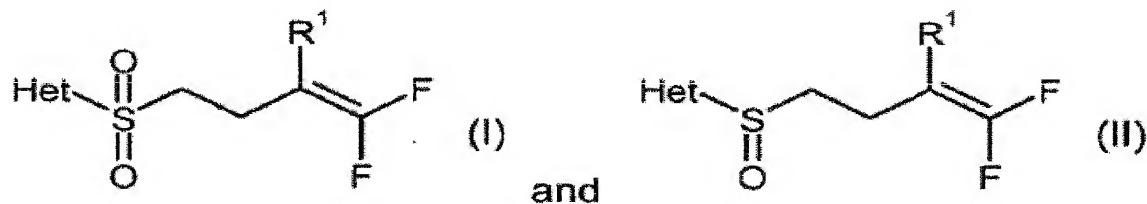
Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

Claim 1-10. (Cancelled)

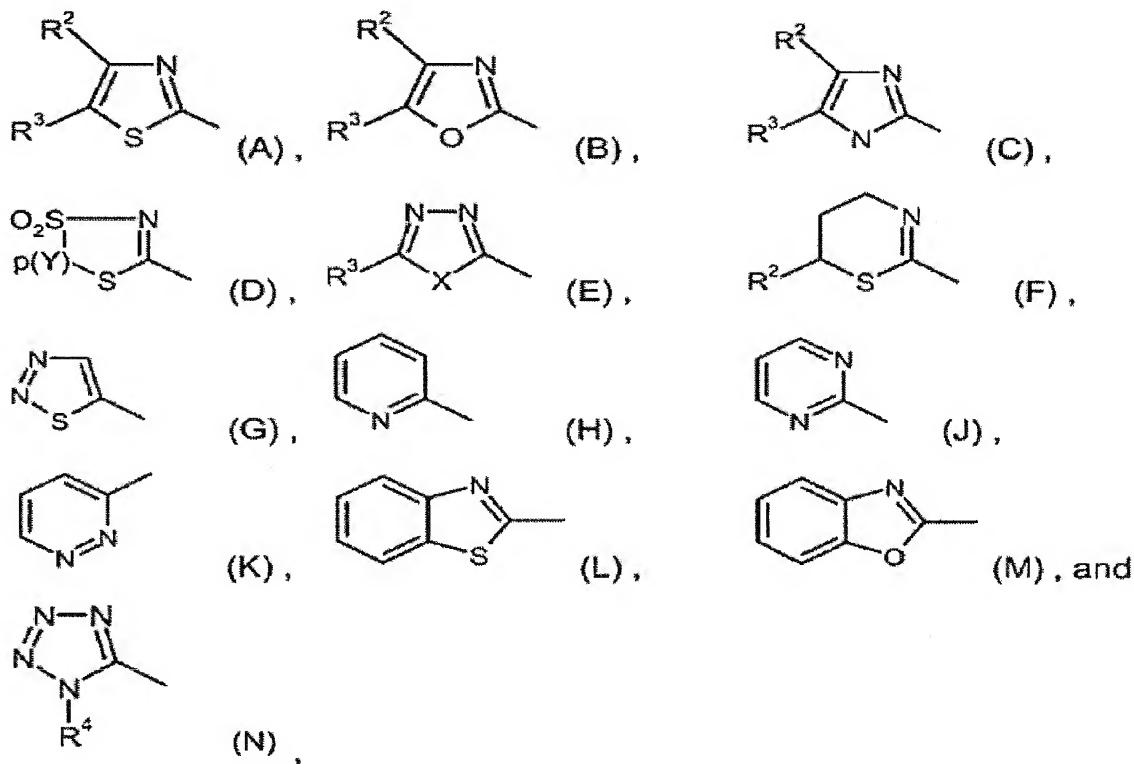
Claim 11 (Previously Presented): A process for preparing heterocyclic fluoroalkenyl sulfone and sulfoxide compounds of formulas (I) and (II)



where

R¹ is hydrogen or fluorine, and

Het is a heterocycle selected from the group consisting of



where

R² is hydrogen, halogen, C₁-C₂-alkyl, or C₁-C₄-haloalkyl,

R³ is hydrogen or halogen; or is optionally halogen-, methyl-, ethyl-, n- or i-propyl-, n-, i-, s-, or t-butyl-, methoxy-, ethoxy-, n- or i-propoxy-, or n-, i-, s-, or t-butoxy-substituted C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-alkylsulfinyl, C₁-C₄-alkylsulfonyl, C₁-C₄-alkoxycarbonyl, C₁-C₄alkoxy-C₁-C₄-alkyl, C₁-C₄-alkylthio-C₁-C₄-alkyl, carboxyl, C₁-C₄-alkylaminocarbonyl, C₃-C₆-cycloalkylaminocarbonyl, C₁-C₄-dialkylaminocarbonyl, C₂-C₄-alkenyl, C₂-C₄-alkenylthio, C₂-C₄-alkenylsulfinyl, or C₂-C₄-alkenylsulfonyl,

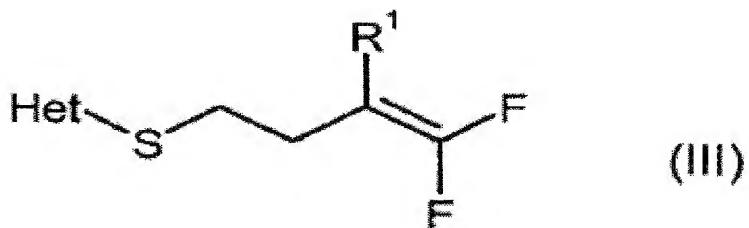
R is C₁-C₈-alkyl, C₂-C₆-alkenyl, C₁-C₄-haloalkyl, C₁-C₄-alkoxy-C₁-C₄-alkyl, C₁-C₄-alkylthio-C₁-C₄-alkyl, or C₃-C₈-cycloalkyl; or is optionally halogen-, C₁-C₄-alkyl-, C₁-C₄-alkoxy-, C₁-C₄-alkylthio-, or C₁-C₄-haloalkyl-substituted phenyl or benzyl,

p is 1, 2, or 3,

X is oxygen or sulfur, and

Y is methylene that is optionally singly or doubly, identically or differently, substituted with optionally halogen-, C₁-C₄-alkoxy-, C₁-C₄-alkylthio-, C₁-C₄-haloalkoxy-, or C₁-C₄-haloalkylthio-substituted C₁-C₄-alkyl, C₂-C₄-alkenyl, or C₂-C₄-alkynyl; or is phenyl that is optionally singly to triply, identically or differently, substituted with halogen, cyano, nitro, C₁-C₄-alkyl, C₁-C₄-alkoxy, C₁-C₄-alkylthio, C₁-C₄-haloalkyl, C₁-C₄ haloalkoxy, or C₁-C₄-haloalkylthio,

comprising allowing a compound of formula (III)



where R¹ and Het are each as defined for formula (I),
to react with a salt of peroxomonosulfuric acid (H₂SO₅),

optionally in the presence of a reaction assistant and optionally in the presence of a diluent, wherein the reaction of a compound of formula (II) to formula (I) is conducted at a pH of from 6 to 10.

Claim 12. (Cancelled)

Claim 13. (Cancelled)

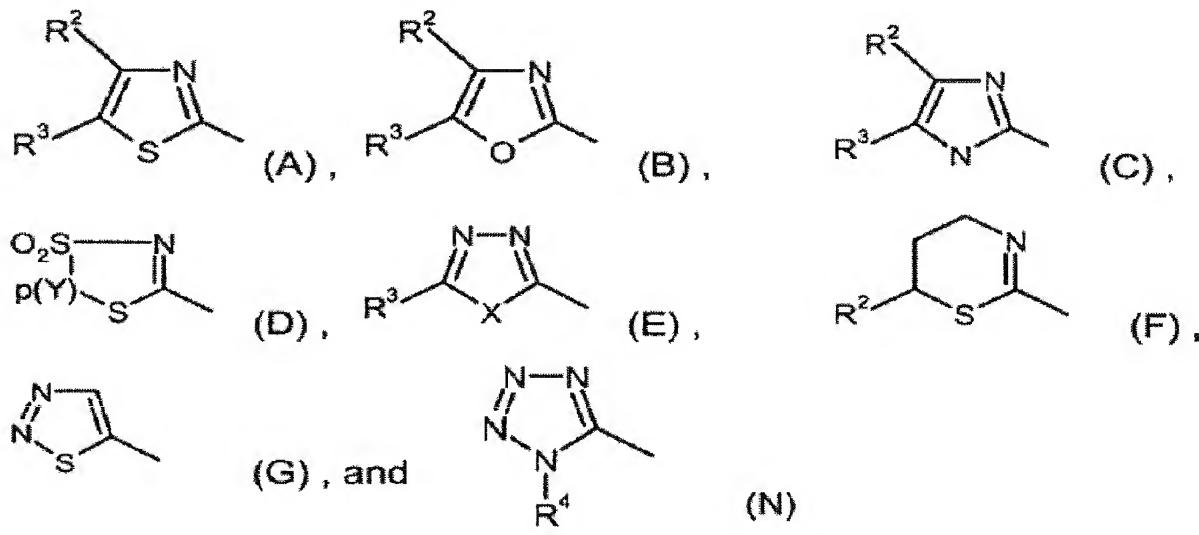
Claim 14. (Previously Presented) A process for preparing compounds of formula (II) according to Claim 11 wherein a compound of formula (III) according to Claim 11 is allowed to react with a salt of peroxomonosulfuric acid (H_2SO_5), optionally in the presence of a reaction assistant and optionally in the presence of a diluent.

Claim 15. (Previously Presented) A process according to Claim 14 carried out at a pH of from 1 to 3.

Claim 16. (Previously Presented) A process according to Claim 11 in which the salt of peroxomonosulfuric acid is potassium hydrogenperoxomonosulfate ($2 KHSO_5 \bullet KHSO_4 \bullet K_2SO_4$ (5:3:2:2)).

Claim 17. (Previously Presented) A process according to Claim 11 carried out at a temperature of from -20°C to 150°C.

Claim 18. (Previously Presented) : A process according to Claim 11 in which R¹ is fluorine, Het is a heterocycle selected from the group consisting of

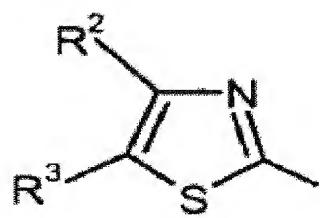


R² is hydrogen, fluorine, or chlorine,
R³ is hydrogen, fluorine, or chlorine; or is optionally fluorine-, chlorine-, methyl-, ethyl-, n- or i-propyl-, n-, i-, S-, or t-butyl-, methoxy-, ethoxy-, n- or i-propoxy-, n-, i-, S-, or t-butoxy-substituted methyl, ethyl, n- or i-propyl, n-, i-, S-, or t-butyl, methoxy, ethoxy, n- or i-propoxy, n-, i-, S-, or t-butoxy, methylthio, ethylthio, nor i-propylthio, n-, i-, S-, or t-butylthio, methylsulfinyl, ethylsulfinyl, methylsulfonyl, ethylsulfonyl, methoxycarbonyl, ethoxycarbonyl, n- or i-propoxycarbonyl, n-, i-, S-, or

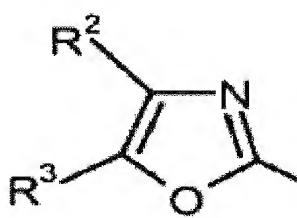
t-butoxycarbonyl, methoxymethyl, methoxyethyl,
ethoxymethyl, ethoxyethyl, methylthiomethyl,
methylthioethyl, ethylthiomethyl, ethylthioethyl,
carboxyl, methylaminocarbonyl, ethylaminocarbonyl, n- or
i-propylaminocarbonyl, cyclopropylaminocarbonyl,
cyclobutylaminocarbonyl, cyclopentylaminocarbonyl,
cyclohexylaminocarbonyl, dimethylaminocarbonyl,
diethylaminocarbonyl, ethenyl, propenyl, or butenyl, R₄ is
methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl,
tert-butyl, n-pentyl, cyclopropyl, cyclopentyl,
cyclohexyl, 2-chloroethyl, 2,2,3,3,3-pentafluoropropyl,
2,2,2-trifluoroethyl, 3-bromopropyl, 2-methoxyethyl, 2-
ethoxyethyl, 2-methylthioethyl, allyl, or 2-butenyl; or
is optionally singly or doubly, identically or
differently, fluorine-, chlorine-, bromine-, methyl-,
ethyl-, isopropyl-, trifluoromethyl-, methoxy-, or
methylthio-substituted phenyl or benzyl,
P is 1 or 2,
X is oxygen, and
Y is methylene that is optionally singly or doubly,
identically or differently, substituted with methyl or
ethyl; or is phenyl that is optionally singly to triply,
identically or differently, substituted with fluorine,

chlorine, methyl, methoxy, trifluoromethyl, cyano, or nitro.

Claim 19 (Previously Presented): A process according to Claim 11 in which Het is a heterocycle selected from the group consisting of



(A) and

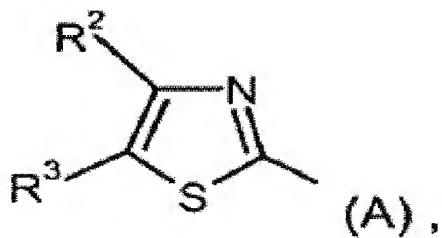


(B),

R² is hydrogen, and

R³ is hydrogen, fluorine, or chlorine.

Claim 20 (Previously Presented): A process according to Claim 11 in which



R² is hydrogen, and

R³ is chlorine.

21. (New) A process for preparing a compound of formula (I) as defined in Claim 11, wherein a compound of formula (II) as defined in claim 11 is allowed to react with a salt of peroxomonosulfuric acid (H_2SO_5), optionally in the presence of a reaction assistant and optionally in the presence of a diluent, wherein the process is conducted at a pH of from 6 to 10.